\$/079/62/032/005/008/009 D204/D307

5.3630 AUTHORS:

Moshkina, T.M., and Pudovik, A.N.

TITLE:

Synthesis of glycol diphosphates and of certain deri-

vatives of phosphinic acids

PERIODICAL: Zhurnal obshchey khimii, v. 32, no. 5, 1962, 1671-1675

TEXT: A series of diphosphates of ethylene, diethylene and tetraethylene glycols, 1,4-butanediol, β-thiodiglycol, N-methyl diethanolamine and nitropropylene glycol was synthesized, owing to the potential application of such compounds as plasticizers. Two methods
tential application of such compounds as plasticizers. Two methods
were used, giving 25 - 75 % yields: (1) Dialkyl (or diaryl) phosphowere used, giving 25 - 75 % yields: (1) Dialkyl (or diaryl) phosphoric chloroanhydrides were added dropwise to an ethereal solution of
ric chloroanhydrides were added dropwise to an ethereal solution of
the appropriate glycol, in the presence of pyridine, at 0-5 °C. The
mixture was stirred for a further 1 hr. at 25 - 30°C. Pyridine hydrochloride was filtered off, the filtrate washed with water, which
was then frozen out, and the ether was removed by distillation.(2)
Phosphorus oxychloride was added to cooled glycols (0 - 5°C) and
Phosphorus oxychloride was added to cooled glycols (0 - 5°C) and
the mixture was stirred for 1 hr., removing the HCl formed. The resulting dichloroanhydride was added dropwise to the appropriate alCard 1/2

PUDOVIK, A.N.; ALADZHEVA, I.M.

Polyphosphites. Part 2: Reactions of dialkyl phosphorous chlorides with aromatic dioxy compounds. Zhur.ob.khim. 32 nc.6:2005-2010 Je 162. (MIRA 15:6)

(Diphosphites) (Aromatic compounds)

MUKNENEVA, N.A.; KIRPICHNIKOV, P.A.; PUDOVIK, A.N.

Polyphosphites. Part 3: Interaction of pyrocatesholphosphorous chloroanhydride with dioxy compounds. Zhur.ob.khim. 32 no.7:2193-2196 Jl 162. (MIRA 15:7)

1. Kazanskiy khimiko-tekhnologicheskiy institut imeni S.M.Kirova. (Phosphorous acid) (Pyrocateshol) (Diphosphites)

PUDOVIK, A.N.; KONOVALOVA, I.V.

Regrouping of methyldi-(diethylphosphone)-carbinol. Dokl.
AN SSSR 143 no.4:875-878 Ap '62. (MIRA 15:3)

1. Kazanskiy gosudarstvennyy universitet im. V.I.Ul'yanova-Lenina. Predstavleno akademikom B.A.Arbuzovym. (Phosphinic acid) (Phosphorus acids)

/145/002/011/018 B106/B101

15 8110

AUTHORS:

Pudovik, A. N., Yevstaf'yev, G. I., and Cherkasov, R. A.

TITLE:

Addition of incomplete phosphorus acid esters to unsaturated

polyesters

Akademiya nauk SSSR. Doklady, v. 145, no. 2, 1962, 344-346 PERIODICAL:

TEXT: This is a continuation of previous papers on the addition of various phosphorus acid esters to unsaturated electrophilic reagents in the presence of basic catalysts. Polyesters with molecular weights between 700 and 4000 obtained by condensation of maleic anhydride with various glycols were made to react with various esters of phosphorous acid. The resulting phosphorus-containing polyesters may be of interest as plasticizers, and as a component for the production of refractory material. Excessive addition of diethyl phosphite to polyethylene glycol maleinate (molecular weight: 750) in the presence of little sodium methylate as a catalyst, is a very vigorous and exothermic reaction yielding a solid, hygroscopic resin which does not continue burning when taken out of flame. Diethyl phosphite is added to all polyester double bonds. Experiments

Card 1/3

Addition of incomplete phosphorus ...

S/020/62/145/002/011/018 B1C6/B101

with mixtures of diethyl phosphite and polydiethylene glycol maleinate (molecular weight: 3600) of different molar ratios showed that the diethyl phosphite is added practically quantitatively to the polyester double bonds. Adequate choice of molar ratios allows the production of polyesters containing any desired amount of phosphonic groups and double bonds. Polydiethylene glycol maleinate was also made to react with diethyl thiophosohite and two cyclic alkyl phosphites (phosphorous ester with diethylene glycol, or 1,3-butylene glycol). Phosphite was always used in excess to guarantee the addition to all polyester double bonds. Furthermore, the reaction of diethyl dithiophosphate with the condensation products of maleic annydride and ethylene glycol,  $\beta$ -thiodiglycol, 1,4-butylene glycol, and diethylene glycol (molecular weights of the polyesters: 1053-3183) was studied at different molar ratios. The products of addition to all polyester double bonds were rubber-like or solid materials, e.g., of the structure HO[-CO-CH-CH\_-COOCH\_CH\_O-]\_H. The products  $s = P(OC_2H_5)_2$ 

of incomplete addition were viscous resins. All these polyesters are

Card 2/3

## "APPROVED FOR RELEASE: 06/15/2000

## CIA-RDP86-00513R001343530001-1

Addition of incomplete phosphorus ...

S/020/62/145/002/011/018 B106/B101

insoluble in alcohol and dioxane, some dissolve in water and yield opalescing solutions. Polyesters continue burning when taken out of flame owing to their sulfur content. Successive addition of diethyl phosphite and diethyl dithiophosphate to some unsaturated polyesters was also conducted. The ratios were chosen in such a way that the two phosphorus compounds added to 50% of the polyester double bonds. The polyesters thus obtained are viscous, water-soluble resins or solids which continue burning when taken out of flame. There are 4 tables.

ASSOCIATION: Kazanskiy gosudarstvennyy universitet im. V. I. Ul'yanova-Lenina (Kazan' State University imeni V. J. Ul'yanov-Lenin)

PRESENTED:

January 23, 1962, by B. A. Arbuzov, Academician

SUBMITTED:

January 16, 1962

Card 3/3

PUDOVIK, A.N.; KASHEVAROVA, E.I.; KHUSAINOVA, N.G.

Phosphorus-containing amides of acrylic and methacrylic acids.

Dokl.AN SSSR 145 no.4:818-821 Ag 162. (MIRA 15:7)

1. Kazanskiy gosudarstvennyy universitet im. V.I.Ul'yanova-Lenina. Predstavleno akademikom B.A.Arbuzovym.

(Acrylamide) (Methacrylamide) (Phosphorus organic compounds)

<u>L 12425-63</u>

EWP(j)/EPF(c)/EWT(m)/EDS

ASD Pc-4/Pr-4 RM/WW S/0190/63/005/006/0886/0891

66

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ACCESSION NR: AP3001162

AUTHOR: Pudovik, A. N.; Yevstaf'yev, G. I.

TITLE: Synthesis of phosphorus-containing polyesters by the polytransesterification reaction

SOURCE: Vy\*sokomolekulyarny\*ye soyedineniya, v. 5, no. 6, 1963, 886-891

TOPIC TAGS: polyesters, transesterification, synthesis, glycols, diethyl ethylphosphinite, diethylphosphite

ABSTRACT: The objective of the present investigation was the study of polytransesterification of glycols with diethyl-ethylphosphinite and diethylphosphine. The reagents were used in equimolecular amounts, the reaction being controlled by alcohol yield, at 175, 180, 190, and 200C. It was found that the reaction rate and degree of its completion goes up with the temperature. The molecular weights of the obtained polyesters, as determined by the cryoscopic method, showed little differences. In the reaction with 1,4-butanediol the formation of tetrahydrofurane was observed along with the polyester, which suggests the cyclization of 1,4-butanediol. A 65-70% yield of cyclic acids was obtained in 30 minutes at 130C and a 100 mm pressure. Orig. art. has: 4 formulas, 5 charts, and 1 table.

Association: Kazan State University Cord 1/2/

L 14945-63 EWP(j)/EPF(c)/EWT(m)/BDS ASD Pc-4/Pr-4 RM/WW

ACCESSION NR: AP3003799 S/0190/63/005/007/1106/1110

AUTHORS: Moshkins, T. M.; Pudovik, A. N.

TIME: Polyethyleneglycols and their derivatives

64

SOURCE: Vy\*sokomolekulyarny\*ye soyedineniya, v. 5, no. 7, 1963, 1106-1110

TOPIC TAGS: polyethyleneglycol, ethylene oxide polymer, ethylene glycol, mono-

ABSTRACT: Polymerization of ethylene oxide was conducted in flasks containing 0.08 Mol ethyleneglycol, 30 ml benzene, and 0.0008 Mol boron trifluoride etherate, through which ethylene oxide was bubbled at 40-450 for a period of 15-17 hours. The obtained polymers were waxy white compounds. These were fractionated by fractional precipitation with ethyl ether from 25 benzene solutions. The polymerization coefficients of the fractions, everaging 40-60, were determined from viscosimetric measurements in dioxane solution by Ostwald's method. The synthesis of polyethyleneglycol-monochloroacetates was achieved by slowly adding to polyethyleneglycol at OC an equimolar quantity of chloroacetic acid, the resulting products representing highly viscous fluids or vaseline-like masses, soluble in ethanol, benzene, dioxane and carbon tetrachloride. When used as a catalyst in the polymerization of

Card 1/2

ethylene oxide, products with a 50-70 polymerization coefficient were obtained. A still higher polymerization coefficient of 65-85 was recorded when diethyl phosphate replaced boron trifluoride etherate in a similar setup. Orig. art. has: 2 formulas and 2 cherts.			
ASSOCIATION: Kazanskiy filial nauchno-issledovatel'skogo kinofoto-institut (Kazan Division of Scientific Research, Kinophoto Institute)			
SUBMITTED: 17Jan62	DATE ACQ: 08Aug63	<b>1</b> 2	ICL: 00
UB CODE: CH	NO REF SOV: 004	Оті	R: 007
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PUDOVIK, A.N.; KHUSAINOVA, N.G.; KASHEVAROVA, E.I.

Polymerization and copolymerization of phosphorus-containing esters of methacrylic acid. Vysokom.soed. 5 no.9:1376-1381 S '63.

(MIRA 17:1)

1. Kazanskiy gosudarstvennyy universitet imeni V.I.Ul'yanova-Lenina.

PUDOVIK, A. N.; MOSHKINA, T. M.; KHRAMTSOVA, V. P.

Diazophosphinic and hydrazodiphosphinic esters. Zhur. ob. khim. 33 no.1:94-97 '63. (MIRA 16:1)

1. Kazanskiy filial Nauchno-issledovatel'skogo kinofotoinstituta.

(Phosphinic acid) (Diazo compounds)
(Hydraso compounds)

PUDOVIK, A. N.; KONOVALOVA, I. V.

Interaction of carboxyl chloride and carboxylic anhydrides with sodium diethyl phosphite. Zhur. ob. khim. 33 no.1: 98-102 163. (MIRA 16:1)

1. Kazanskiy gosudarstvennyy universitet.

(Acids, Organic) (Phosphorous acid)

PUDOVIK, A. N.; ALADZHEVA, I. M.; SOKOLOVA, I. A.; KOZLOVA, G. A.

Polyphosphites. Part 4: Reactions of dialkyl phosphoryl chlorides with glycols. Zhur. ob. khim. 33 no.1:102-107 '63. (MIRA 16:1)

1. Kazanskiy gosudarstvennyy universitet.

(Phosphoryl chloride) (Glycols)

PUDOVIK, A.N.; KONOVALOVA, I.V.; DEDOVA, L.V.

Rearrangement of esters of hydroxymethyl (diethylphosphone) acetic acid. Zhur.ob.khim. 32 no.2:483-486 F '63.

(MIRA 16:2)

1. Kazanskiy gosudarstvennyy universitet.

(Acetic acid) (Rearrangements (Chemistry))

PUDOVIK, A.N.; ALADZHEVA, I.M.

Acetylene-allene-acetylene rearrrangements of phosphites with a 3, /-acetylene bond in an ester radical. Zhur.ob.khim. 33 no.2:707-7-8 F '63. (MIRA 16:2)

1. Kazanskiy gosudarstvennyy universitet. (Phosphorous acid) (Rearragmements (Chemistry)) (Phosphinic acid)

PUDOVIK, A.N.; ALADZHEVA, I.M.

Acetylene-allene-diene rearrangements fo diphosphites with a  $\beta$ ,  $\gamma$  - acetylene bond in a common ester radical. Zhur.ob.khim. 33 no.22708-709 F '63. (MIRA 16:2)

1. Kazanskiy gosudarstvemnyy universitet.
(Diphosphorous acid) (Rearrangements (Chemistry))
(Butadiene)

PUDOVIK, A.N.; KHUSAINOVA, N.G.; ALADZHEVA, I.M.

Reaction of nucleophilic addition to alkynyl phosphinites. Zhur.ob.khim. 33 no.3:1045-1046 Mr '63. (MIRA 16:3)

1. Kazanskiy gosudarstvennyy universitet.
(Phosphinic acid) (Unsaturated compounds)
(Addition reactions)

PUDOVIK, A.N.; KRUPNOV, G.P.

Reactions of nucleophilic addition of phosphoric acid dialkyl esters amides. Zhur. ob. khim. 33 no.5:1654-1658 My '63. (MIRA 16:6)

1. Kazanskiy filial Vsesoyuznogo nauchno-issledovatel'skogo kinofotoinstituta.

(Phosphoric acid)
(Addition reactions)

L 9907-63 EWP(j)/EFF(c)/EWP(q)/EWT(m)/

BDS\_AFFTC/ASD\_Pr-4/Pc-4-RM/MAY/WW/JD

s/0079/63/033/006/1816/1821

ACCESSION NR: AP3002623

AUTHOR: Pudovik, A. N.; Aladzheva, I. M.

TITLE: Polyphosphites. V. Synthesis and properties of cyclic diphosphites

SOURCE: Zhurnal obshchey khimii, v. 33, no. 6, 1963, 1816-1821

TOPIC TAGS: cyclic diphosphites, phosphorous acid, phosphorous acid esters, 1,3-ethenediol, 1,2-propanediol, 1,3-propanediol, 1,3-butanediol, bis(2-hydroxyethyl) ether, pyrocatechol, polyphosphites, polymers

ABSTRACT: The synthesis and properties of certain cyclic diphosphites having an aliphatic chain or an aromatic ring as the common radical have been studied for the first time. The diphosphites were synthesized in a 50 to 70% yield from the cyclic phosphorochloridites and various glycols or pyrocatechol, with cooling in absolute ethyl ether in the presence of triethylemine. The diphosphites are liquids readily soluble in many organic solvents, react exothermically with water,

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L 9907-63

AP3002623 ACCESSION NR:

Some of their physical constants and are stable in storage and distillation. and yields are given in table 1 of the Enclosure. The diphosphites enter all reactions characteristic of P(III)-containing compounds; triethylene diphosphite, for example, reacts with an equimolar amount of sulfur to form triethylene bisthiophosphate. Reactions of the phosphites with haloalkanes are of general interest. Thus, triethylene diphosphite and bromoethane undergo an Arbuzov rearrangement at 1300 to form ethylene ethylphosphonate and dibromoethane, probably by mechanism B as shown in Fig. 1 of the Enclosure. Arbuzov rearrangement can be used to prepare polymers from the diphosphites. 7 Heating of a diphosphite with an equimolar amount of a dihaloalkane such as dibromoethane or dibromobutane produces highly viscous, resinous P-containing polymers. Heating of a diphosphite with a catalytic quantity of iodoethane produces hard, glassy polymers which are slightly soluble in organic solvents. The preparation and properties of the polymers will be described in greater detail in a separate paper. Orig. art. has: 3 formulas and 1 table.

ASSOCIATION: Kazanskiy gosudarstvenny\*y universitet (Kazan' State University) DATE ACQ: 20Jul63 ENCL: SUBMITTED: 21May62

SUB CODE:

00

NO REF SOV: 009

OTHER: 001

Card

PUDOVIK, A.N.; KONOVALOVA, I.V.; ISHMAYEVA, E.A.

Reactions of the diene synthesis and addition of butadienephosphinic and butadienethiophosphinic esters. Zhur. ob. khim. 33 no.8: 2509-2513 Ag '63. (MIRA 16:11)

1. Kazanskiy gosudarstvennyy universitet.

PUDOVIK, A.N.; TARASOVA, R.I.; BULGAKOVA, R.A.

Reactions of sodium diethyl thiophosphite with haloallyl compounds. Zhur. ob. khim. 33 no.8:2560-2563 Ag '63.

(MIRA 16:11)

1. Kazanskiy gosudarstvennyy universitet.

PUDOVIK, A.N.; KUZOVLEVA, R.G.

Reactions of nucleiophilic reagents with vinylphosphinic and acetoxyvinylphosphinic acid esters. Zhur. ob. khim. 33 no.8:2755-2760 Ag '63. (MIRA 16:11)

1. Kazanskiy gosudarstvennyy universitet.

MUKHENEVA, N.A.; KIRPICHNIKOV, P.A.; PUDOVIK, A.N.

Polyphosphites. Part 6: Interaction of diaryl phosphoryl chlorides with dihydroxy compounds. Zhur.ob.khim. 33 no.10:3192-3196 0 '63. (MIRA 16:11)

PUDOVIK, A.N.; MURATOVA, A.A.; SEMKINA, E.B.

Reactions of dialkylphosphinic acid esters with trialkyl tin halides. Zhur.ob.khim. 33 no.10:3350-3353 0 '63. (MIRA 16:11)

1. Kazanskiy gosudarstvennyy universitet.

PUDOVIK, A.N.; PUDOVIK, M.A.

New method of synthesizing phosphinic and thiophosphinic acid esters. Part 40: Addition of acid phosphites, diphosphites, phenylphosphine to unsaturated electrophilic reagents containing one or two double bonds. Zhur.ob.khim. 33 no.10: 3353-3358 0 163. (MIRA 16:11)

1. Kazanskiy gosudarstvennyy universitet.

PUDOVIK, A.N.; GAREYEV, R.D.

Reactions of carbethoxycarbene with unsaturated organophosphorus compounds. Zhur.ob.khim. 33 no.10:3441-3442 0 '63. (MIRA 16:11)

1. Kazanskiy gosudarstvennyy universitet.

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PUDOVIK, A.N.; KONOVALOVA, I.V.

Transformations of allylphosphinic esters in the presence of sodium ethylate. Zhur.ob.khim. 33 no.10:3442-3443 0 '63. (MIRA 16:11)

1. Kazanskiy gosudarstvennyy universitet.

PUDOVIK, A.N.; ALADZHEVA, I.M.; YAKOVENKO, L.N.

Synthesis and rearrangement of aiethylpropargyl phosphite. Zhur.ob.khim. 33 no.10:3443-3444 0 163. (MIRA 16:11)

1. Kazanskiy gosudarstvennyy universitet.

是我们是我们的一个人,我们也是我们们的一个人,我们们也不是我们的,我们们们们的一个人,我们们们们们的一个人,我们们们们的一个人,我们们们们们的一个人,我们们们们

PUDOVIK, A.N.: KONOVALOVA, I.V.

Synthesis of styrene and its homologs by pyrolysis of phosphates. Dokl. AN SSSR 149 no.5:1091-1094 Ap 163. (MIRA 16:5)

l. Kazanskiy gosudarstvennyy universitet im. V.I.Ul yanova-Lenina. Predstavleno akademikom B.A.Arbuzovym.

(Styrene) (Phosphates)

PUDOVIK, A.N.; ALADZHEVA, I.M.

Thermal or "pseudoclaisen rearrangements of allyl and propargyl esters of phosphorous acid. Dokl. AN SSSR 151 no.5:1110-1113 Ag '63. (MIRA 16:9)

1. Kazanskiy gosudarstvennyy universitet im. V.I.Ul'yanova-Lenina.
Predstavleno akademikom B.A.Arbuzovym.
(Phosphorous acid) (Esters) (Rearrangements (Chemistry))

ACCESSION NR: AP4017637

5/0190/64/006/002/0253/0264

AUTHORS: Pudovik, A. N.; Muratova, A. A.; Sushentsova, F. F.; Zoreva, N. M.

TITLE: Heterochain polymers with phosphorus and oxygen atoms in the main chain. Polyphosphinophosphates and polyphosphinates

SOURCE: Vy\*sokomolekulyarny\*ye soyedineniya, v. 6, no. 2, 1964, 258-264

TOPIC TAGS: polymer, polycondensation, phosphinic acid, alkylphosphinic acid, alkylphosphinic acid ester, alkylphosphinyl dichloride, phosphoryl dichloride, ethyldichlorophosphine, polyphosphinophosphate, polyphosphinate, heterochain polymer

ABSTRACT: This investigation involved polyphosphinophosphates (PPP) and polyphosphinates (PP), the polymeric chain of which consisted of links

$$\begin{bmatrix}
 OR & R' \\
 -P - O - P - O - P - O \\
 0 & O
\end{bmatrix}$$
and
$$\begin{bmatrix}
 R & R' \\
 -P - O - P - O \\
 0 & O
\end{bmatrix}$$

with radicals containing from 2 to 11 carbons. These polymers were obtained by

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ACCESSION NR: AP4017637

polycondensation of alkylphosphinic acid esters with dichlorides of alkylphosphoric-, alkylphosphinic-, and arylphosphinic acids. The polycondensation was conducted for 4-10 hours at a gradual temperature rise from 120 to 200C. The molecular weight, softening point, and solubility of the obtained polymers in water and in organic solvents were determined. It was found that the PPP compounds, which contained 4-8 carbon atoms per link, dissolved only in water and alcohols and were insoluble in organic solvents. An increase in the number of carbon atoms to 14 per link resulted in the formation of polymers soluble in organic solvents, possessing a low melting point from -30 to -50C, displaying good adhesion to glass, and having a low flammability. The replacement of an aliphatic radical by benzyl raised the melting point by about 60-80C. The PPP and PP compounds are rapidly hydrolyzed by water (even at OC). When the molecular ratio of the issuing alkylphosphinic acid esters and of the dichlorides was 1:1, the polymerization yielded only products of low molecular weight (676-712). A 30% excess of dichloride was required to bring it up to 2600-2890. It was found that the investigated polymerization reactions were of the second order, and that the reaction rate increased with temperature, as well as in the presence of such catalysts as FeCla ZnCl2, and AlCl3. Orig. art. has: 2 charts, 4 formulas, and 3 tables.

Card 2/3

ACCESSION NR: AP4017637

ASSOCIATION: Kazanskiy gosudarstvenny\*y universitet im. V. I. Lenina (Kazan'

SUB CODE: CH

State University)

DATE ACQ: 23Mar64

ENCL: 00

SUBMITTED: OlDec62

NO REF SOV: 003

OTHER: 003

PUDOVIK, A.N.; TARASOVA, R.I.

Reactions of di- and triarylhalomethanes with salts of phosphinic, phosphorothioic, and phosphorodithioic acids. Zhur.ob.khim. 34 no.1: 293-298 Ja 64. (MIRA 17:3)

1. Kazanskiy gosudarstvennyy universitet.

PUDOVIK, A.N .; FAYZULLIN, E.M.; ZHURAVLEV, G.I.

Mechanism and order of addition of phosphorus trichloride and other phosphoryl chlorides to propylene exide. Dokl. AN SSSR 165 no.3:586-589 N '65. (MIRA 18:11)

1. Kazanskiy gosudarstvennyy universitet im. V.I. Ul'yanova-Lenina. 2. Chlen-korrespondent AN SSSR (for Pudovik).

ACC NR: AP7003661  SOURCE CODE: UR/0079/66/036/008/1454/1459  ACTIOR: AP7003661  SOURCE CODE: UR/0079/66/036/008/1454/1459  ACTIOR: AP7003661  SOURCE CODE: UR/0079/66/036/008/1454/1459  CRG: Kazan' State University im. V. I. Ul'yanov-Lenin (Pazanskiy Zosudarstvennyy universitet)  TITLE: Reactions of olefin oxides with phosphorus oxychloride and diethyl chlorophosphate  SCURCE: Zhurnal obshchey khimii v. 36, no. 8, 1966, 1454-1459  SCURCE: Zhurnal obshchey khimii v. 36, no. 8, 1966, 1454-1459  SCURCE: Zhurnal obshchey khimii v. 36, no. 8, 1966, 1454-1459  SCURCE: Zhurnal obshchey khimii v. 36, no. 8, 1966, 1454-1459  SCURCE: Zhurnal obshchey khimii v. 36, no. 8, 1966, 1454-1459  SCURCE: Code in the presence of small quantities  ADSTANCT: It was found that phosphorus oxychloride and diethyl chlorophosphate  ADSTANCT: It was found that phosphorus oxychloride actid. The presence of small quantities  are readily added to alpha-exides of olefins win the presence of small quantities  are readily added to alpha-exides of traces of water or hydrochloric actid.  did not proceed at all in the absence of traces of water or hydrochloric actid.  did not proceed at all in the absence of traces of water or hydrochloric actid.  did not proceed at all in the absence of traces of water or hydrochloric actid.  A reaction of cheminary carbon atom. Sproposed. In the reaction of  the converted to a glycol chlorohydrin, is proposed. In the reaction of  the side of the primary carbon atom. When the olefin oxides are treated with  on the side of the primary carbon atom. When the olefin oxides are treated with  phosphorus oxychloride with propylene oxide, opening of the oxide ring occurs  phosphorus oxychloride, monochlorides, dichlorides, and complete esters  amount of hydrogen chloride, monochlorides, dichlorides, and complete esters  amount of hydrogen chloride, monochlorides, dichlorides, and complete esters  of the corresponding beta-chloroalkylphosphoric acids are obtained. A series  of the corresponding beta-chloroalkylphosphoric	<u> Paragonal Bergeratus de Caralles de la companya d</u>	শ্বীকর্মনুর
universitet) TITLE: Reactions of olefin oxides with phosphorus oxychloride and diethyl chlorophosphate SOURCE: Zhurnal obshchey khimil v. 36, no. 8, 1966, 1454-1459 SOURCE: Zhurnal obshche v. 36, 1966, 1454-1459 SOURCE: 241, 14 SOURCE: Zhurnal obshche v. 36, 1966, 1454-1459 SOURCE: Zhurnal obshche v. 36,	ACC MAI MITOUST 1	
	universitet) TITLE: Reactions of olefin oxides with phosphorus oxychloride and diethyl chlorophosphate SCURCE: Zhurnal obshchey khimii v. 36, no. 8, 1966, 1454-1459 TOPIC TAGS: ethylene oxide, organic oxide, organic phosphorus compound TOPIC TAGS: ethylene oxide, organic oxide, organic phosphorus compound ABSTRACT: It was found that phosphorus oxychloride and diethyl chlorophosphate are readily added to alpha-oxides of olefins in the presence of small quantities are readily added to alpha-oxides of olefins oxides tested were ethylene oxide, of water or hydrochloric acid. The olefin oxides tested were ethylene oxide, of water or hydrochloric acid. The olefin oxides tested were ethylene oxide, of water or hydrochloric acid. did not proceed at all in the absence of traces of water or hydrochloric acid. A reaction scheme involving the formation of an oxonium intermediate, which is a reaction scheme involving the formation of an oxonium intermediate, which is then converted to a glycol chlorohydrin, is proposed. In the reaction of then converted to a glycol chlorohydrin, is proposed. In the reaction of then converted to a glycol chlorohydrin, when the olefin oxides are treated with on the side of the primary carbon atom. When the olefin oxides are treated with on the side of the primary carbon atom. When the olefin oxides are treated with on the side of the primary carbon atom. When the olefin oxides are obtained. A series of the corresponding beta-chloroalkylphosphoric acids are obtained. A series of the corresponding beta-chloroalkylphosphoric acids were obtained by the of dialkyl-beta-chloroalkyl esters of phosphoric acids with alcohols.  Together the tables.  TOPIC 1000 TOPIC 1	

1. T1805-0; 1.WC(M)/EMF(J) (M) ACC NR: AP7003662	SOURCE CODE: UR/0079/66/036/008/1460/1467
AUTHOR: Pudovik, A. M.; Durova, O. O.G.: Institute of Organic Chemistr khimil AN SSSR)  TITLE: Addition of incomplete esterologically and butynone  SOURCE: Churnal obshchey khimil, v. TOPIC TAGS: ester, organic phosphorous acid are added to the simplest represendance and ketones propargyl of alkali metal alcoholates at the hydroxyphosphinic acids. This reach alpha, beta-unsaturated ketones. It is also bond, yielding mixed esters of dithest type of unsaturated organophosphorous and extenses of dithest type of unsaturated organophosphorous alleres.	rs of phosphorus acids to propargyl  36, no. 8, 1966, 1460-1467  Trus compound, mercaptan lete esters of phosphorus acids: dialkyl- acids and acid esters of ethylphosphinous esentatives of alpha, beta-acetylenic aldehyde and butynone in the presence carbonyl group, forming esters of acetylenic tion was in contrast to that of ethylenic bialkyl esters of dithiophosphoric acid react copargyl aldehyde at the carbon-carbon triple diophosphoric acid. Representatives of a new as compounds were synthesized: diphosphonal- a was found to react readily with ethyl muct. Orig. art. has: 1 figure and 1 table.
Card 1/1 jb	UDC: 547.341

L 11h06-67 EUT(m)/EWT(j) RM/JW SOURCE CODE: UR/0079/66/036/008/1467/1472 ACC NR: AP7003663 AUTHOR: Pudovik, A. N.; Pudovik, M. A. ORG: Institute of Organic Chemistry, Kazan' (Institut organicheskoy khimii) TITLE: Atomic refraction of phosphorus in esters of alkylarylphosphinic acids, arylphosphinous acids, and reactions of addition to unsaturated compounds SOURCE: Zhurmal obshchey khimii, v. 36, no. 8, 1966, 1467-1472 TOPIC TAGS: phosphinic acid, ester, acrylic acid, methacrylic acid ABSTRACT: A series of eight acid esters of phonylphosphinous acid and eight acid esters of p-tolylphosphinous acid were synthesized. The atomic refraction of the pentavalent phosphorus atom in esters of phenylphosphinous acid was found to be 5.60, and in esters of p-tolyiphosphinous acid 6.10. The atomic refraction of the pentavalent phosphorus atom in esters of alkylphenylphosphinic acids has a value of 5.3h. Replacement of the alkonyl group by a phonyl in dialkylphosphorous and esters of alkylphosphinic acids leads to an increase in the atomic refraction of pentavalent phosphorus to 1.08. The addition of acid esters of p-tolylphosphorous acid to unsaturated electrophilic compounds: esters of acrylic and methacrylic acids, acrylonitrile, and Schiff's bases, were studied. In the presence of sodium alcoholate, the reaction proceeded exothermically, with 52-60% yields of the addition products. Monoesters of tolylphosphinous acid and acid phosphites were also added to unsaturated electrophilic reagents containing two double bonds: dibenzal-p-phenylenediamine and diamis produced from terephthalic aldehydes, aniline and p-chloroanaline. Orig. art. has: 5 tables. [JPRS: 38,970] SUB CODE: 07 / SUBM DATE: 12Jul65 / ORIG REF: 004 / GTH REF: 001 UDC: 547,26'118 Card 1/1 jb

UR/0020/66/166/003/0615/0618 EWT(m)/EMP(j) L 35386-66 SOURCE CODE: AP6026818 ACC NR: AUTHOR: Pudovik, A. N. (Corresponding member AN SSSR); Gazizov, T. Kh.; Samitov, Yu. Yu.; Zykova, T. V. ORG: Institute of Organic Chemistry, AN SSSR, Kazan' (Institut organicheskoy khimii AN SSSR) TITIE: Reaction of dialkyl acetyl phosphites with chloral SOURCE: AN SSSR. Doklady, v. 166, no. 3, 1966, 615-618 TOPIC TAGS: phosphorus compound, chemical composition, chemical bonding, IR spectrum ABSTRACT: The authors studied the reaction between dialkyl acetyl phosphites and chloral. Acetyl chloride was not observed in the products of reactions of dimethyl-, diethyl- and di-n-propylacetylphosphates with chloral with a yield of 70-75%. An analysis of these products shows that they correspond to the composition CC13CHO·(RO)2POCOCH3. There is no adsorption in the infrared spectrum in the 1680-1620 cm<sup>-1</sup> region which is characteristic for valency vibrations of the double carbon-carbon bond. There are bands which are characteristics for the P=0 bond in the 1280 cm-1 region and for P=0-R groups in the 1070-1020 cm-1 region. Orig. art. has: 1 figure and 2 tables. SUB CODE: 07, 20 / SUBM DATE: 09Jul65 / ORIG REF: 008 / OTH REF: 005 546.183.315+547.446.1

EWT(m)/EWP(j) 34130-66 SOURCE CODE: UR/0079/66/036/001/0161/0162 ACC NR: AP6025539 AUTHOR: Pudovik, A. N.; Ishmayeva, E. A.; Akhmorova, R. S.; Aladzheva, I. M. ORG: Kazan! Stato University im. V. I. Ul!yanov-Lenin (Kazanskiy gosudarstvennyy universitet) TITLE: Addition of mucleophilic reagents to 2,3-di(diethylphosphone)-butadiene-1,3/ SOURCE: Zhurnal obshchey khimii, v. 36, no. 1, 1966, 161-162 TOPIC TAGS: phosphorus acid, phosphorus compound, exothermic reaction, IR spectrum, potassium compound ABSTRACT: The addition of nucleophilic reagents: dimethyl- and diethylphosphorous acids, ethyl mercaptan, and diethylamine to butadiene was found to proceed in the presence of alcoholates of the alkali metals. Addition proceeds exothermally in the 1,2-position. l-Dimethylphosphone-2,3-di(diothylphosphone)butene-3, 1,2,3-tri(diethylphosphone)butene-3, 1-mercaptocthyl-2,3-di(diethylphosphone)butene-3, and 1-diethylamino-2,3-di(diethylphosphone)butene-3 were synthesized; their structures were established by study of their infrared spectra and oxidation with potassium permanganate. JPAS: 35,9987 SUB CODE: 07. 20 / SUBN DATE: 21Apr65 / ORIGREF: 001 547.26 1118

ACC NR. AP7000240 SOURCE CODE: UR/0079/66/036/004/0718/0724 AUTHOR: Pudovik, A. N.; Fayzullin, E. M.; Zhuravlev, G. I. ORG: Kazan' State University im. V. I. Ul'yanov-Lenin (Kazanskiy gosudarstvennyy universitet) TITLE: Reactions of alpha-oxides with dialkyldithiophosphoric and dithiophosphinic Moscow. Zhurnal Obshchey Khimii, Vol 36. No 4, 1966, pp 718-724 Abstract: The addition of dialkyldithiophosphoric and diphenyldit hiophosphinic acids to nonsymmetrical alpha-oxides or olefins, e.g. glycerin epichlorohydrin, propylene, divinyl, styrene, and glycidol oxides, was studied. The reactions proceed readily without catalysts and are accompanied by a substantial thermal effect. Conclusions on the structure of the addition products and the order of addition of dithioacids to alpha-oxides (in accord with the Markovnikov rule) were drawn on the basis of a study of the chemical properties and infrared spectra of the products. The acid esters of dithiophosphoric and diphenyldithiophosphinic acids were found to be electrophilic in reactions with alpha-oxides, the reactions proceeding with preliminary formation of intermediate oxonium compounds. In the reaction of ethyleneglycol chlorohydrin, 2,3-propyleneglycol chlorohydrin, and

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	iichlorohydrin wi ling ethyl esters has: 1 table.	th the potassium salt of 0,S-alkyleneglycol /JPRS: 37,177	of diethyldithiopho dithiophosphoric ac	osphoric acid, the cid were obtained.	
TOPIC TAGS	: IR spectrum,	organic phosphorus com	pound, ester		
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	05163-67 EWT(m)/EWP(j)/EWP(t)/ETI IJP(c) JD/RM SOURCE CODE: UR/0062/66/000/006/1123/1124
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í	SHAGIDULLIN, R. R., CHERNOVA, A. V., ISHMAYEVA, E. A., PUDOVIK, A. N., Institute of Organic and Physical Chemistry imeni A. E. Arbuzov, Academy of Institute of Organic high correspondency i fizicheskoy khimii AN SSSR)
	Institute of Organic and Physical Chemistry Institute Organic And Physical Chemistry In
į	Sciences USSR (Institute of guille and a the Thompson Atom
-	Question of Conjugation with Participation of the Phosphorus Atom"
	room graden Vhimicheskava, No 6, 1966,
	Moscow, Izvestiya Akademii Nauk SSSR, Seriya Khimicheskaya, No 6, 1966,
	nn 1102-1124
	channel charmtion spectra and Raman spectra of
	compounds containing a diene group with substituents -r OR (OR)
	investigated in comparison with isoprene. In the case of a terminal situation
	investigated in comparison with isoprene. In the case of a serimum in the of the substituents, a bathochromic shift of the absorption maximum in the of the substituents, a bathochromic shift of the absorption maximum in the
	of the substituents, a dathornionic state the intensity of the bands in
	ultraviolet spectrum and a sharp increase in the intensity of the valence vibra- the infrared spectrum and lines in the Raman spectrum of the valence vibra- the infrared spectrum and lines in the Raman spectrum of the valence vibra-
	the infrared spectrum and lines in the Raman spectrum of the value of the c=C bonds is observed. These signs of conjugation are more protions of the C=C bonds is observed. In the case of side substitution by
	nounced for the carbonyl substitution
	two -P groups, the opposite picture is observed: hypsochromic shift in
	two -P groups, the opposite picture is observed: hypsochromic shift in
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	the ultraviolet spectrum, decrease in intensities in the infrared and Ram spectra, indicating a decrease in the conjugation between the two C=C bone probably due to competition by the phosphorus portions of the molecule. I liminary results on the intensities of the bands of the P=O and P=S bonds indicate the participation of these bonds in conjugation. Orig. art. has:  2 formulas.	ds, Pre→	
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L 04095-67 EWP(j)/EWT(l)/EWT(m)/T IJP(c) RM
ACC NR: AR6023276 SOURCE CODE: UR/0058/66/000/003/D124/D124
THOR: Pudovik, A. N.; Moshkina, T. M.; Krupnov, G. P.; Bukin, A. I.; Semenova, L. A.
THE: Plastification of triacetate celluloid films by mixed phosphoric-acid ethers
URCE: Ref zh. Fizika, Abs. 3D1028
URCE: Ref zh. Fizika, Abs. 3D1028
PIC TAGS: photographic film, plasticizer
STRACT: The authors investigated the plastification of triacetate films by mixing cosphoric-acid ethers. It is shown that at least some diphosphates of diethylene ycol result in better mechanical film properties than the previously used mixture triphenyl phosphate and dibutyl phthalate. However, in the presence in them of iphatic radicals, their compatibility with the film deteriorates with increasing angth of the radical. To improve the compatibility, one can introduce cyclic radicals, atoms, and alcoxyl groups into the ether groups. The most effective for the comtibility are the latter, and they also improve noticeably the physical and mechanical operties of the films. A. Karuzhanskiy. [Translation of abstract]
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3h021-66 www(m)/EWF(g) SOURCH CODE: UR/0079/66/036/001/0069/0073 ACC NR: A16025532 AUTHOR: Pudovik, A. H.; Klasainova, N. G.; Galoyova, R. G. B QiG: Kazan' State University (Kazanskiy gosudarstvennyy universitet) TITLE: Addition of compounds with a labile hydrogen atom in the methylene group to esters of propynylphosphinic acid 1 Zhurnal obshchey khimii, v. 36, no. 1, 1966, 69-73 TOPIC TAGS: hydrogen atom reaction, malonic ester, ester, chemical bonding, tautomorism ABSTRACT: Compounds of labile hydrogen atoms in the methylene group: malonic. cyanoacetic, acetoacetic, and phosphonoacetic esters and their homologs are added in the presence of sodium alcoholate to dialkyl esters of propymylphosphinic acid to form addition products at the triple bond. There is no further addition of substances with active methylene groups at the double bonds of the addition products under the experimental conditions selected. The high observed values of the molecular refractions of the addition products and the absence of exaltation of the molecular refraction indicate the presence of keto-enol tautomerism, with predominately the enol form. The addition of selenophenol to the diethyl ester of propynylphosphinic acid gave the mono addition product 24% yield. Orig. art. has: 1 table. [JPRS: / SUBM DATE: 22Sep64 / ORIG REF: 002 / OTH REF: 003 Card 1/1 *574 •468: 547 • 393* 

SOURCE CODE: UR/0079/66/036/003/0494/0498 ACC NR: APPOZZZÓ83 AUTHOR: Khayrullin, V. K.; Pudovik, A. N. OdG: Institute of Organic Chemistry, AN SSSR, Kazan' (Institut organicheskoy khimii AN SSSR) TITLE: Reaction of ethyldichlorophosphine with crotonic acid B SOURCE: Zhurnal obshchoy khimii, v. 36, no. 3, 1966, 494-498 TOPIC TAGS: alkylphosphine, chlorinated organic compound, esterification, IR spectrum, chemical synthesis ABSTRACT: The reaction of ethyldichlorophosphine with crotonic acid was found to proceed smoothly at room temperature, forming a single product, 3-methyl-4-oxo-4-chloro-4-phosphacaproyl chloride. Reaction of 3-methyl-4-oxo-4-chloro-4-phosphacaproyl chloride with alcohols yielded esters of 3\_methyl\_4\_oxo\_4\_alkoxy\_4\_phosphacaproic acid. In these reactions, anlydrides of ethyl-beta-carbalkoxyisopropylphosphinic acid are formed at the same time. Infrared spectra are cited for the chloride and five esters. Orig. art. has: l figure and 2 tables. JPRS7 SUB CODE: 07 / SUBM DATE: 05Apr65 / ORIG REF: 002 Card 1/1 15 547.26 118

L 31799-66 EWT(m)/EWP(1) WW/RM SOURCE CODE: UR/0079/66/036/003/0565/0565

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AUTHOR: Pudovik, A. N.; Pudovik, M. A.

61

ORG: Institute of Organic and Physical Chemistry, AN SSSR, Kazan' (Institut organichoskoy i fizichoskoy khimil AN SSSR)

TITLE: Addition of acid cyclic diothylono glycol diphosphite at multiple bonds

SOURCE: Zhurnal obshchey khimii, v. 36, no. 3, 1966, 565

TOPIC TAGS: reaction mechanism, chemical bonding, organic phosphorus compound, heterocyclic base compound, chemical reaction

ABSTRACT: Acid cyclic diethylene glycol diphosphite is capable of undergoing reactions of nucleophilic addition to unsaturated electrophilic compounds containing C=C, C=O, and C=W bonds. Phonyl and naphthyl isocyanates, acrylonitrile, p-dimethyl-aminobenzaldehyde, p-bromobenzaldehyde, and benzalaniline were used as the unsaturated compounds. The reaction was carried out at 100° with an excess of the unsaturated compound in the presence of alkali metal alcoholates. It can be carried out in the absence of a solvent or in anhydrous alcohol solution. Orig. art. has: 1 table.

[JPRS]

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IDC: 547.26 118

-EWP(j)/EWT(m UR/0079/65/035/008/1502/1503 SOURCE CODE: ACC NR AP6019329 Pudovik, A. N.; Krupnov, G. P. AUTHOR: ORG: Kasan' State University (Kasanskiy gosudarstvennyy universitet) TITIE: Addition of diethylphosphorous acid anilide to Schiff bases SOURCE: Zhurnal obshohey khimii, v. 35, no. 8, 1965, 1502-1503 TOPIC TAGS: Schiff base, organic phosphorus compound, amide, chemical reaction By the addition of diethylphosphorous acid anilide to Schiff bases (I) substances (II) were prepared: (EtO)2PNHPh+PhCH=NC6H4R (I) - (EtO)2P(=NPh)-CHPh-NHC6H4R (II) R = H, p-Me, p-NO<sub>2</sub> The structure of (II) was confirmed by the fact that sulfur did not add to them on heating to  $140^{\circ}$  and by the formation of aniline hydrochloride from (II) (R = H) on hydrolysis in acidified water. Furthermore, trialkyl phospites do not add to Schiff bases in a spontaneous reaction of the type by which compounds (II) formed i.e., the amide group of diethylphosphorous acid anilide must have participated in the reaction. Compounds (II) were crystalline Cord 1/2

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	Eb m. n. 17	2-3°, 165-6°,	and 187-90	for (II) (E	= H),	
(TT) (R =	p-Me) and	(II) (R = p-	NC2), resp.	Orig. art. ha	s: 3 formulas	•
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L 31275-66 EWT(m)/EWP(1) RM SOURCE CODE: UR/0079/66/036/002/0296/0302 ACC NR: AP6022799	
72	٠.
AUTHOR: Khayrullin, V. K.; Sobchuk, T. I.; Pudovik, A. N.	
ORG: Institute of Organic Chemistry, AN SSSR, Kazan' (Institut organicheskoy khimii	
AN SSSR)	
TITLE: Reaction of ethyldichlorophosphine with alpha, beta-unsaturated acids	
SOURCE: Zhurnal obshchey khimii, v. 36, no. 2, 1966, 296-302	
TOPIC TAGS: reaction mechanism, chlorinated organic compound, IR spectrum, esterification, chemical synthesis, alkylphosphine, diene synthesis	
ABSTRACT: The addition of ethyldichlorophosphine to propiolic acid proceeds in the 1,4-position with the formation of 4-oxo-4-chloro-4-phospha-2-hexenoyl chloride. In the reaction of 4-oxo-4-chloro-4-phospha-2-hexenoyl chloride with alcohols and diethyl-amine, esters and the diethylamide of 4-oxo-4-alkoxy-4-phospha-2-hexenoic acid were obtained. Esters of 4-oxo-4-alkoxy-4-phospha-2-hexenoic acid, containing an electrophilic bond, readily add nucleophilic reagents and are extremely active in diene synthesis. The nucleophilic reagent is directed to the beta-carbon atom with	
products. The authors thank R. R. Shagidullin for producing and interpreting the IR spectrum. Orig. art. has: 1 figure and 1 table. [JPRS]	
SUB CODE: 07 / SUBM DATE: 15Mar65 / ORIG REF: 006 / OTH REF: 002	
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AUTHOR: Pudovik, A. N.; Fayzullin, E. M.; Zhukov, V. P.  ORG: Kazan' State University (Kazanskiy gosudarstvennyy universitet)  TITLE: Cyclic esters of unsaturated phosphinic acids  SOURCE: Zhurnal obshchey khimii, v. 36, no. 2, 1966, 310-314  TOPIC TAGS: esterification, cyclic group, organic phosphorus compound, chlorinated organic compound, isomerization, molecular structure, chemical decomposition, phosphinic acid  ABSTRACT: A series of alkyleneglycol-beta-chloroalkyl estors of phosphorus acid were prepared in high yields by the action of chlorides of alkylenegly-colphosphorous acids on ethylene oxide, propylene oxide, and glycerol epichlorohydrin. The cyclic esters of phosphorous acid added sulfur when heated to 100-110°, being converted to esters of thiophosphoric acid. Thermal isomerization (180-200°) of alkyleneglycol-beta-chloroalkyl esters of phosphorous acid yielded alkyleneglycol esters of beta-chloroalkylphosphinic acids. An Arbuzov rearrangement also occurred upon heating of the cyclic phosphities with alkyl halides. Treatment of the alkyleneglycol esters of beta-chloroalkylphosphinic acids with triethylamine in benzene solution with heating resulted in splitting off hydrogen chloride, and formation of alkyleneglycol esters of vinyl- and propenylphosphinic acids. Orig. art. has:  3 tables. [JPRS]  SUB CODE: 07 / SUEM DATE: OSMar65 / ORIG REF: CO5	L 31270-66 EWT(m)/EWF(j) RM
AUTHOR: Pudovik, A. N.; Fayzullin, E. M.; Zhukov, V. P.  ORC: Kazan' State University (Kazanskiy gosudarstvennyy universitet)  TITLE: Cyclic esters of unsaturated phosphinic acids  SOURCE: Zhurnal obshchey khimii, v. 36, no. 2, 1966, 310-314  TOPIC TAGS: esterification, cyclic group, organic phosphorus compound, chlorinated organic compound, isomerization, molecular structure, chemical decomposition, phosphinic acid  ABSTRACT: A sories of alkyleneglycol-beta-chloroalkyl esters of phosphorus acid were prepared in high yields by the action of chlorides of alkylenegly-colphosphorous acids on ethylene oxide, propylene oxide, and glycerol epichlorohydrin. The cyclic esters of phosphorous acid added sulfur when heated to 100-110°, being converted to esters of thiophosphoric acid. Thermal isomerization (180-200°) of alkyleneglycol-beta-chloroalkylphosphinic acids. An Arbuzov rearrangement also occurred upon heating of the cyclic phosphites with alkyl halides. Treatment of the alkyleneglycol esters of beta-chloroalkylphosphinic acids with triethylemine in benzene solution with heating resulted in splitting off hydrogen chloride, and formation of alkyleneglycol esters of vinyl- and propenylphosphinic acids. Orig. art. has:  SUB CODE: 07 / SUBM DATE: OSMar65 / ORIG REF: CO5	
	AUTHOR: Pudovik, A. N.; Fayzullin, E. M.; Zhukov, V. P.  ORG: Kazan' State University (Kazanskiy gosudarstvennyy universitet)  TITLE: Cyclic esters of unsaturated phosphinic acids  SOURCE: Zhurnal obshchey khimii, v. 36, no. 2, 1966, 310-314  TOPIC TAGS: esterification, cyclic group, organic phosphorus compound, chlorinated organic compound, isomerization, molecular structure, chemical decomposition, phosphinic acid  ABSTRACT: A series of alkyleneglycol-beta-chloroalkyl esters of phosphorus acid were prepared in high yields by the action of chlorides of alkylenegly-colphosphorous acids on ethylene oxide, propylene oxide, and glycerol epichloronydrin. The cyclic esters of phosphorous acid added sulfur when heated to 100-110°, being converted to esters of thiophosphoric acid. Thermal isomerization (180-200°) of alkyleneglycol-beta-chloroalkyl esters of phosphorous acid yielded alkyleneglycol esters of beta-chloroalkylphosphinic acids. An Arbuzov rearrangement also occurred upon heating of the cyclic phosphites with alkyl halides. Treatment of the alkyleneglycol esters of beta-chloroalkylphosphinic acids with triethylamine in benzene solution with heating resulted in splitting off hydrogen chloride, and formation of alkyleneglycol esters of vinyl- and propenylphosphinic acids. Orig. art. has:  3 tables. [JPRS]
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#### CIA-RDP86-00513R001343530001-1

L 27714-66- EWT(m)/EWP(j) ACC NR. AP6018514 SOURCE CODE: UR/0079/65/035/011/2080/2081 AUTHOR: Pudovik, A. N.; Ishmayeva, E. A. ORG: none TITLE: Addition of S-chlorodiethylthiophosphate to diviny SOURCE: Zhurnal obshchey khimii, v. 35, no. 11, 1965, 2080-2081 TOPIC TAGS: IR spectrum, hydrolysis, oxidation, organic phosphorus compound, organic sulfur compound, chlorinated organic compound ABSTRACT: The reaction of S-chlorodiethylthiophosphate with the simplest representative of dienic compounds with a conjugated system of double bonds -divinyl -- at -50 results in the production of a small amount of the diethyl ester of monothiophosphoric acid and an addition product in 60% yield. The infrared spectrum of the addition product and results of its oxidation with potassium permanganate and hydrolysis indicated that addition of S-chloroethylthiophosphate to divinyl proceeds in the 1,2-position, yielding (EtO) P (=0)-S-CH2-CHC1-CH=CH2. The formation of a small amount of the diethyl ester of monothiophosphoric acid is explained by partial hydrolysis of the S-chlorodiethylthiophosphate during the reaction and treatment of the reaction mixture. Orig. art. has: 1 formula. /JPRS/ SUB CODE: 07/ SUBM DATE: 19Apr65/ ORIG REF: 002/ OTH REF: 005 UDC: 547.26 118/547.368

 $L_26577-66$  EWT(m)/EPF(n)-2/EWP(j) RM/JD,

ACC NR. AP6016977

SOURCE CODE: UR/0020/65/165/003/0586/0589

AUTHOR: Pudovik, A. N. (Corresponding member AN SSSR); Fayzullin, E. M.; Zhuravlev, G. I.

40 B

ORG: Kazan' State University im. V. I. Ul'yanov-Lenin (Kazanskiy gosudarstvennyy universitet).

TITLE: Mechanism and order of addition of phosphorus trichloride and other chlorides of phosphorus acids to propylene oxide of phosphorus acids to propylene oxide

SOURCE: AN SSSR. Doklady, v. 165, no. 3, 1965, 586-589

TOPIC TAGS: phosphorus chloride, ester, tertiary amine, hydrolysis, IR spectrum, phosphorous acid, hydrogen chloride

ABSTRACT: New evidence confirming the proposed mechanism of the reactions of phosphorus trichloride and chlorides of incomplete esters of phosphorus acids with alpha-olefins (through preliminary opening of the oxide ring by hydrogen chloride) was obtained in an investigation of the reaction of propylene oxide with phosphorus trichloride and the chloride of dibutylphosphorus acid. The reactions proceeded readily when the reagents, were combined. However, when a small amount of triethylemine was added to the reaction mixtures, these reactions did not take place. If anhydrous propylene oxide was added to the dibutylphosphorous acid chloride, freshly distilled under vacuum, and protected from moisture, no reaction between them was observed. The introduction of

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ACC NR: AP6016977

atmospheric air containing moisture into the reaction volume of the addition of one to two drops of water to the reaction mixture gave rise to a vigorous reaction. The authors conclude that the first step in the reactions considered is a partial hydrolysis of the acid chlorides and interaction of the hydrogen. chloride thereby formed with the alpha-oxide. The oxonium ion formed upon addition of a proton to the oxide then either directly reacts with the chloride ion, to form propylene glycol chlorohydrin, which then reacts with the scid chloride, or perhaps simultaneously with the chloride ion, molecules of the chlorohydrin are involved in the reaction. When the chlorine atoms are replaced by alkoxyl groups, the basicity of the phosphorus atom decreases, which facilitates the reactions of the acid chloride with the oxide. A mixed ester of phosphorous acid is formed, and hydrogen chloride is regenerated. The reaction of propylene oxide with phosphorus trichloride was conducted in ether solution with cooling, at ratios of 1:1, 2:1, and 3:1; the dichloride of betachloroisopropylphosphorous acid, and tri-beta-chloroisopropyl phosphite were obtained in good yields. The presence of a secondary alcohol group in the investigated chlorohydrin was corroborated by the infrared spectra and chemical investigations. Orig. art. has: 1 figure and 1 table. [JPRS]

SUB CODE: 07 / SUBM DATE: 10Mar65 / ORIG REF: 004 / OTH REF: 002

Card 2/2

#### "APPROVED FOR RELEASE: 06/15/2000

#### CIA-RDP86-00513R001343530001-1

ENT(m)/T/EWP(j)/ETC(m)=6 L 25978-66 WM/RM IJP(a) ACC NR: AP6015614 SOURCE CODE: UR/0020/66/168/002/0354/0356 AUTHOR: Pudovik, A. N.; Pudovik, M. A. ORG: Institute of Organic and Physical Chemistry im. A. Ye. Arbuzov AS SSSR, Kazan (Institut organicheskoy i fizicheskoy khimii Akademii nauk SSSR) TITLE: Migration polymerization of acid diphosphites, diphosphinites [sic] with p-tolylene diisocyanate and some other compounds with two double bonds SOURCE: AN SSSR. Doklady, v. 168, no. 2, 1966, 354-356 TOPIC TACS: phosphorus containing polymers, fire resistant material, migration polymerization, para tolylene diisocyanate, acid phosphite ABSTRACT: Migration polymerization of acid diphosphites with p-tolylene diisocyanate (I) or with some other compounds was studied. Cyclodiethylene glycol diphosphite (II) or diethyl 1,3-propylene glycol diphosphite (III) [designated "diphosphinite" in the original] were used as phosphorus-containing components. The effects of temperature, polymerization time, ratio of components and the nature and amount of solvents used in some experiments as polymerization media on the yields and properties of the polymers obtained were studied. Most of the experiments were conducted with components in the equimolecular ratios, in the absence of solvent and in nitrogen atmosphere. In some cases, however, dimethylformamide, dioxane or ethyl acetate

**Card** 1/3

UDC: 546.183:547.315.3

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ACC NR. AP6015614

$$\begin{array}{c} CH_{3} \\ CH_{4} \\ CH_{3} \\ CH_{4} \\ CH_{5} \\ CH_{5$$

were used as media. In the first series of the experiments, I and II produced solid orange colored polymers. With an increase in polymerization time the yields and molecular weight of the polymers also increased until the maximum yield of 78.8% was attained. The softening temperatures also increased until they reached the 120 C range at the above-mentioned maximum yield. Maximum yields were obtained at the equimolar ratio of components. Polymerization in solvent decreased the yields and the molecular weight of the polymers. The effect is based on the solubility of polymers in the given solvent to a definite molecular weight. Polymers obtained from solvents are more uniform and less intensely colored than polymers obtained without solvent. Similar relationships were observed for the polymers obtained from I and

Card 2/3

ACC NR: AP6015614 III. The polymers were yellow or orange powders, soluble only in dimethylformamide or dimethylsulfoxide. Their softening temperatures were above 100 C. Compound III was subjected to migration polymerization with some other compounds which contained two double bonds, such as ethylene glycol dimethacrylate, dibenzal acetone, some Schiff's bases or terephthalic aldehyde. The polymerization was conducted in the absence of solvent, at the equimolar ratios of the components and in the presence of sodium ethylate. Polymers were reprecipitated from dimethylformamide and dried to constant weight. They were either solid powders or viscous resins, soluble in dimethylformamide, dimethylsulfoxide or H2SO4. All of them had a low combustibility, and in some cases, were almost incombustible. Orig. art. has: 1 formula, 2 tables Programme Street [BN] SUB CODE: 07, 11/ SUBM DATE: 30Sep65/ ORIG REF: 003/ OTH REF: ATD PRESS: 005/ Card 3/3 FW

ACC NR: AP6018509

SOURCE CODE: UR/0079/65/035/011/2042/2046

AUTHOR: Moshkina, T. M.; Pudovik, ApN.

,

ORG: none

TITLE: Phosphorus-containing azo-\and hydrazo-compounds

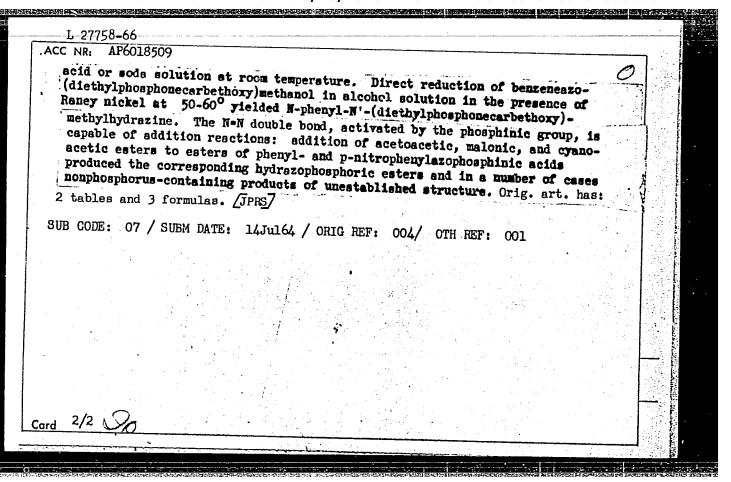
SOURCE: Zhurnal obshchey khimii, v. 35, no. 11, 1965, 2042-2046

TOPIC TAGS: organic azo compound, organic synthetic process, organic phosphorus compound, organic nitrile compound, hydrazine derivative, ester

ABSTRACT: The azo combination of aromatic diazo compounds with a number of organophosphorus compounds containing an activated methylene group: phosphone-acetic ester, phosphoneacetone, and phosphoneacetonitrile was carried out under mild conditions in the absence of catalysts. Some properties of the phosphorus-containing azo-compounds synthesized were studied. They decompose gradually during storage with an evolution of nitrogen; the decomposition process is substantially accelerated at increased temperature (above 50°), with an evolution of nitrogen and a further resinification of the products formed. The azo-compounds obtained are highly sensitive to the action of acids and alkalis, yielding a vigorous evolution of nitrogen, accompanied by partial decomposition and resinification of the products when treated with dilute hydrochloric

Card 1/2

UDC: 547.467/8



I. 21860-66 EWT(m)/EWP(j) RM AP6012651 SOURCE CODE: UR/0079/65/035/002/0358/0363 ACC NR: AUTHOR: Pudovik, A. N.; Ishmayeva, E. A. Kazan' State University (Kazanskiy gosudarstvennyy universitet) Butadienethio- and selenophosphinic esters TITLE: SOURCE: Zhurnal obshchey khimii, v. 35, no. 2, 1965, 358-363 TOPIC TAGS: ester, phosphoric acid, mercaptan, butadiene ABSTRACT: Methyl, n-propyl, and n-butyl esters of 1, 3-butadienethiophosphinic acid were obtained and a study made of the addition of unsaturated esters of phosphoric acids and ethylmercaptan to these esters. The addition of nucleophilic reagents to butadiene thiophosphinic esters occurs in the 1, 4 position. The acid dichloride of 4-chlorobutene-2-selenophosphonic acid and the dimethyl ester of 1, 3-butadiene selenophosphonic acid were obtained and characterized. Orig. art. has: 1 table. [JPRS] SUB CODE: 07 / SUBM DATE: 16Dec63 / ORIG REF: 003 / OTH REF: 001 UDC: 547.315.2:546.183

PUDOVIK, A.N.; GAZIZOV, T.Kh.; SAMITOV, Ya.Yu.; EYROVA, T.V.

Praction of dialkyladetyl phosphites with chloral. Dokl. AN SSSR 166 no.3:615-618 Ja 466. (MARA 19:1)

1. Institut organicheskoy khimii AN SSCR, Kazan'. 2. Chlen-korresuondent AN SSSR (for Pudevik). Submitted July 9, 1965.

PUDOVIK, A.N., KUZOVLEVA, R.G.

Polymerization and copolymerization of  $\infty$  and A-carbalkoxy-vinyl phosphinates. Vysokom, soed, 7 no.9:1539-1542 S '65.

(MIRA 18:10)

1. Kazanskiy gosudarstvennyy universitet im, V.I. Ul'yanova-Lenina,

ACC NR: AP5028480  AUTHORS: Moshkina, T. M.; Pudovik, A. N.; Krupnov, G. P.; Bukin, A. I.; Semenova,  L. A. 44,55  ORG: none  TITLE: Method for obtaining plasticized ester-cellulose films, for instance,	
ORG: none  53  B  1. A. 44,55  ORG: none	
ORG: none  53  B  1. A. 44,55  ORG: none	
ORG: none  B  1 - Market and contain collingua films. for instance.	
which can altaining placticized ester-cellulose films, for instance,	
TITLE: Method for obtaining plasticized ester-certains in the state of the state cellulose films. Class 39, No. 175646 announced by All-Union Scientific Research Motion Picture Institute (Vsesoyuznyy nauchno-issledovatel'skiy kinofotoinstitut)	
SOURCE: Byulleten' izobreteniy i tovarnykh znakov, no. 20, 1965, 64	
TOPIC TAGS: polymer, plasticizer, plastic compound, plastic material, plastic, film	
ABSTRACT: This Author Certificate presents a method for obtaining ester-cellulose films, for instance, triacetate cellulose films, by introducing esters of polybasic acids into a solution of cellulose triacetate. To increase the variety of plasticizers, esters of phosphonoacetic acid are used as the plasticizing agent.	
SUB CODE: 11/ SUBM DATE: 13Jun64	
UDC: 678.544.43 678.049.13.002.2	<i>}</i>
Louis	

KIRPICHNIKOV, F.A.; MUKMENEVA, N.A.; PUDOVIK, A.N.; KOLYUBAKINA, N.S.

Reaction of phosphorous acid esters with 1,1-diphenylethane hydrogeroxide. Dokl. AN SSSR 164 no.5:1050-1053 0 165.

(MIRA 18:10)

- 1. Kazanskiy khimiko-tekhnologicheskiy institut im. S.M.Kirova. 2. Chlen-korrespondent AN SSSR (for Pudovik).

TO GREEK AT THE CONTROL OF THE STATE OF THE

PUDOVIK, A.N.; MOSHKINA, T.M.; KRUPNOV, G.P.: BUKIN, A.I.; SEMENOVA, L.A.; Prinimali uchastiye: KOSTYUKOVA, L.A., laborant; PETROVA, M.G., laborant; TEMIRBAYEV, A.M., inzh.; FAYZULLIN, A.Yu., inzh.; POLOZOVA, L.P., laborant; NAZAROVSKAYA, G.V., laborant

Synthesis and study of organophosphorus plasticizers for the triacetate film bases. Trudy NIKFI no.46:17-25 62. (MIRA 18:8)

PUDOVIE, A.H.; YEVSTAF'YEV, G.I.

Synthesis of phosphorus-containing polyesters by means of homopolyreesterification reaction. Dokl. AN SSSR 164 (MIRA 18:10) no.6:1331-1334 0 165.

1. Kazanskiy gosudarstvennyy universitet im. V.1.Wl'yanova-Lenina.

2. Chlen-korrespondent AN SESR (for Pudovik).

TTIE: Polycondensation of 4-chlorobut-2-enephosphinyl chloride with dihydroxy ompounds  OURCE: Vysokomolekulyarnyye soyedineniya, v. 7, no. 5, 1965, 808-812  OPIC TAGS: polycondensation, polymer, resin, reaction kinetics, activation nergy  BSTRACT: The investigation is an extension of the work of V. V. Korshak, I. A. ribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Carlorebut-2-enephosphinyl chloride (A) with aliphatic and iromatic dihydroxy compounds was investigated. The condensation was carried out from the velocity and extent of reaction were determined by the quantity of 600. The velocity and extent of reaction were determined by the quantity of 101 evolved. The activity of aliphatic glycols in decreasing order of activity (CI evolved. The activity of aliphatic glycol, diethylene glycol, butylene-1, reglycol, dipropylene glycol, and hexaethylene glycol. The activity of arcmatic glycols diminished in the order: pyrocatechin, diphenylolpropane, hydroquinone.	53035-65 EVP(j)/EWT(m) Pc-14 JA CESSION NR: AP5013053	UR/0190/65/007/005/0808/0812 541.64+678.675
OURCE: Vysokomolekulyarnyye soyedineniya, v. 7, no. 5, 1965, 808-812  OPIC TAGS: polycondensation, polymer, resin, reaction kinetics, activation nergy  BSTRACT: The investigation is an extension of the work of V. V. Korshak, I. A. ribova, and M. A. Andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. A. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. andrayeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Folyribova, and M. andrayeva (Izv. AN SSSR. Otd. khim.		2.3
OURCE: Vysokomolekulyarnyye soyedineniya, v. 7, no. 5, 1965, 808-812  OPIC TAGS: polycondensation, polymer, resin, reaction kinetics, activation nergy  BSTRACT: The investigation is an extension of the work of V. V. Korshak, I. A. Bribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and december of the condensation was carried out condensation of 4-chlore-but-2-enephosphinyl chloride (A) with aliphatic and condensation was carried out condensation was carried out condensation was carried out condensation of the velocity and extent of reaction were determined by the quantity of the condensation of activity of aliphatic glycols in decreasing order of activity (GI evolved. The activity of aliphatic glycol, diethylene glycol, butylene-1, as: 2 propylene-1,2-glycol, ethylene glycol, diethylene glycol, butylene-1,	THORS: Pudovik, A. N.; Ishmayeva, E	
OURCE: Vysokomolekulyarnyye soyedineniya, v. 7, no. 5, 1965, 808-812  OPIC TAGS: polycondepsation, polymer, resin, reaction kinetics, activation nergy  BSTRACT: The investigation is an extension of the work of V. V. Korshak, I. A. ribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was car	TLE: Polycondensation of 4-chlorobu	nt-2-enephosphinyl chloride with dihydroxy
OPIC TAGS: polycondensation, polymer, resin, reaction kinetics, activation nergy  BSTRACT: The investigation is an extension of the work of V. V. Korshak, I. A. ribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Poly-indensation of 4-chlore-but-2-enephosphinyl chloride (A) with aliphatic and condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out commatic dihydroxy compounds was investigated. The condensation was carried out condensation was carried out condensation of activity of activity of aliphatic glycols in decreasing order of activity of condensation was carried out condensation.		7 _ 5 1065 R08-812
BSTRACT: The investigation is an extension of the work of V. V. Korshak, I. A. ribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR. Otd. khim. n., 1957, 631). Polyribova, and M. A. Andreyeva (Izv. AN SSSR.	OURCE: Vysokomolekulyarnyye soyedine	niya, v. /, no. ), 170), ccc
ribova, and M. A. Andreyeva (12v. An about the condensation of 4-chlore-but-2-enephosphinyl chloride (A) with aliphatic and condensation of 4-chlore-but-2-enephosphinyl chloride (A) with aliphatic and condensation was carried out condensation.	nergy	
Gl evolved. The activity of all phases glycol, diethylene glycol, butylene-1, as: 2 propylene-1,2-glycol, ethylene glycol, diethylene glycol, butylene-1,	ribova, and M. A. Andreyeva (12v. Andreyeva (12v. Andreweva ) ondensation of 4-chlore-but-2-enephoromatic dihydroxy compounds was investigated in the compound of the compound	sphinyl chloride (A) with aliphatic and stigated. The condensation was carried out
Lycols diminished in we older. Provide the control of the control	Cl evolved. The activity of all places as: 2 propylene-1,2-glycol, ethylen	e glycol, diethylene glycol, butylene-1,

GCESSION NR: AP5013053	한 사용하다면 소설을 가지면 되었다.	보기 하는 사용, 이번 이번 하고, 사용, 사용, 사용, 사용, 사용, 사용, 사용, 유명, 유명, 유명, 유명, 유명, 유명, 유명, 유명, 유명, 유명
an energy of activation of lasters of A and diethylene glasters with a butadiene radio and 3 graphs.  ASSOCIATION: Kazanskiy gosu	iethylene glycol was found to 7.9 ± 2.3 Kcal/mole. Dehydrod lycol and of A and butylenegl cal at the phosphorus atom. (darstvennyy universitet im. V	Orig. art. has: 1 table
(Kazan State University) SUBMITTED: 04Jul64	ENGL: OO	SUB CODE: MT,GC
NO REF SOV: 005	OTHER: OOL	
3.4 1.7 ( · · · · · · · · · · · · · · · · · ·		

PUDOVIK, A.N.; ALADZHEVA, I.M.; YAKOVENKO, L.N.

Synthesis and rearrangements of propargyl phosphites and allenyl phosphonates. Zhur. ob. khim. 35 no.7:1210-1217 J1 '65. (MIRA 18:8)

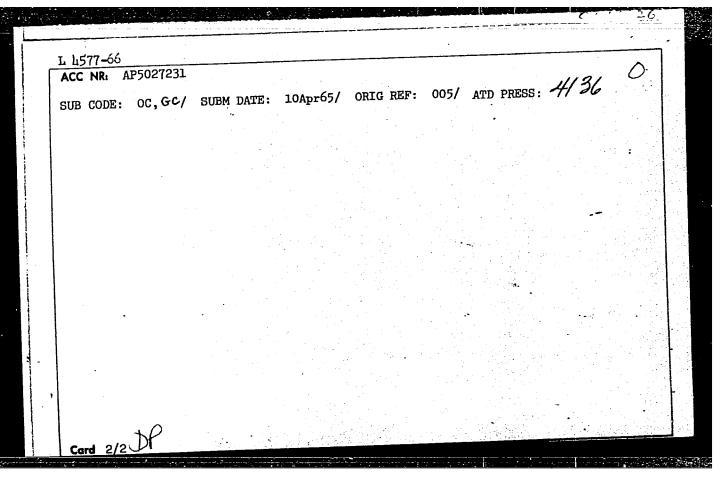
1. Kazanskiy gosudarstvennyy universitet.

L 62796-65 EWT(m)/EPF(c)/EWP(j)/T Pc-li/Pr- ACCESSION NR: AP5018432	66.095.26+678.744
AUTHOR: Pudovik, A. N.; Kashevarova, E. I.; G	oloven'kina, L. I.
TITLE: Polymerization and copolymerization of phates, beta-methacryloxyethyldialkyl phosphit rylic acid and dialkylphosphorous acids	beta-methacryloxyethyldialkyl phos- es and mixed anhydrides of methac-
SOURCE: Vysokomolekulyarnyye soyedineniya, v	7, no. 7, 1965, 1248-1253
TOPIC TAGS: methacrylic acid, organophosphoru	
ABSTRACT: The synthesis of a series of new esphosphorus in the ester radical was carried of phosphates and phosphites studied were obtained dialkyl chlorophosphates and phosphites, as for	ed by reacting ethylene glycol with
$CH_{3}=C (CH_{3}) COOCH_{3}CH_{3}OH + CIP(O)(OR)_{3} \rightarrow CH_{3}=C (CH_{3}) COOCH_{3}CH_{3}OH + CIP(OR)_{3} \rightarrow CH_{3}=C (CH_{3}) COOCH_{3}CH_{3}OH + CIP(OR)_{4} \rightarrow C$	CH <sub>1</sub> =C (CH <sub>2</sub> ) COOCH <sub>2</sub> CH <sub>2</sub> OP (OR) <sub>2</sub> I <sub>2</sub> =C (CH <sub>2</sub> ) COOCH <sub>2</sub> CH <sub>2</sub> OP(OR) <sub>2</sub>
The effect of temperature and nature of the 3	nitiator (0.2 mole % benzoyl peroxide
Cord 1/3	

CESSION NR: AP50	18432					の は 過
0.2 mole % azoiso	butyrodinitrile	e) on the po	olymeriz	ation rat	e of β-metha	cryloxy-
hyldiethyl phospha	te was investig	gated; the r	relation	ships obs	erved also a	bbia to
methacryloxyethylothylothylothyl methacrylate	was studied at	70° C in th	ie prese	nce of 0.	5 mole % ben	zovl per-
ide. To prepare m	ixed anhydrides	of methaci	ylic ac	id and ac	id esters of	phos-
orous acids, the m	eaction of phos	phorous aci	ld anhyd	lrides wit	h potassium	methacry-
te was employed, a	is follows:					
	O I JOR	CH <sub>3</sub>	0	CH.		
	O OR + KOO	OC—C=CH <sub>2</sub> → C	H.POCO—	C≕CH2		
			ÓR			
	o S	ņu,	0	CII		
	(RO,) P CI + KOOC	-C=CH <sub>3</sub> → (RC	), P OCO-	-C=CH <sub>1</sub>		
e anhydrides were		And the second of the second o	and the second of the second		ring vacuum	distilla-
on:						
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	(RO), PCI + KOOC-C=CH			7, 23	
	→ (RO),POCO—C=CH <sub>2</sub> +F	KCl			
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he presence of benzoyl prodinitrile. Mixed and colid polymers insoluble	peroxide and even more in	phorous and methaci	rylic acid form		
the presence of benzoyl payrodinitrile. Mixed and solid polymers insoluble tables.  ASSOCIATION: Kazanskiy	peroxide and even more in nydrides of diethylphosp in organic solvents. (	phorous and methac Orig. art. has: 3	rylic acid form figures and 4		
the presence of benzoyl production trile. Mixed and solid polymers insoluble tables.	peroxide and even more in	phorous and methac Orig. art. has: 3	cylic acid form figures and 4 yanova-Lenina		
the presence of benzoyl payrodinitrile. Mixed and solid polymers insoluble tables.  ASSOCIATION: Kazanskiy	peroxide and even more in nydrides of diethylphosp in organic solvents. (	phorous and methaci Orig. art. has: 3	rylic acid form figures and 4		
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the presence of benzoyl prodinitrile. Mixed and solid polymers insoluble ables.  ASSOCIATION: Kazanskiy (Kazan State University)  SUBMITTED: 13Aug64	peroxide and even more in hydrides of diethylphosis in organic solvents. ( gosudarstvennyy univers.)	phorous and methaconig. art. has: 3  itet im. V. I. Ul'	cylic acid form figures and 4 yanova-Lenina		

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L 4577-66 EWT(m)/EPF(c)/EWP(j)/T RM SOURCE CODE: UR/0020/65/164/006/±331/1334	
1 4577-00 EAT (ATT) SOURCE CODE. 44553	
ACC NR. APOUZIZED 44,55	
ATTHUOR. Pudovik, A. N. (Corresponding member All Doors,	
ACC NR: AP5027231  AUTHOR: Pudovik, A. N. (Corresponding member AN SSSR); Yevstaf'yev, G. I.  ORG: Kazan State University im. V. I. Ul'yanov Lenin (Kazanskiy gosudarstvennyy Universitet)	
ORG: Kazan State University 1m. V. 1. 92	
universitet) 1,44,55	
ORG: Kazan State University im. v. 1	
TITLE: Synthesis of P	
SOURCE: AN SSSR. Doklady, v. 164, no. 6, 1965, 1331-1334	
containing DOLYMEL, Po-V	- 1
SOURCE: AN SSSR. Doklady, v. 164, no. 6, and solver, polyester, transesterification TOPIC TAGS: polymer, phosphorus containing polymer, polyester, transesterification of methylolphosphinic	<b>:</b>
and with the homopolytranses of 122	
esters:  n (C <sub>2</sub> H <sub>2</sub> O)P(O)CH <sub>2</sub> OH - n C <sub>2</sub> H <sub>2</sub> OH - OPCH <sub>2</sub> R - n	
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R (Carajo) (Colonia - R _ n	
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heated in the absence of any catalyst. Floures Reacti	on :
The reaction mixture was heated in the absence of any catalyst. Product yields were the reaction mixture was heated in the absence of any catalyst. Product yields were 50—60%. Rate studies show that the reaction follows second-order kinetics. Reaction 50—60%. Rate studies show that the reaction of activation was found to be the reaction are initially the reaction are initially that were calculated; the energy of activation was found to be	
to how have sometimes an army of accitation and initials	73
The reaction was found to be 50-60%. Rate studies show that the reaction of activation was found to be reaction for the constants were calculated; the energy of activation was found to be rate constants were calculated; the energy of activation was found to be rate constants were calculated; the energy of activation was found to be rate constants were calculated; the energy of activation was found to be rate constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated; the energy of activation was found to be rate constants.  [Violation of the constants were calculated with the constants were calculated with the constants were calculated with the constants.  [Violation of the constants were calculated with the constants were calculated with the constants were calculated with the constant was also constants.]	5]
E = 25,400 I recover liquids with mol. wt = 700-800. On run sate has: 3 figures.	
E = 23,400 ± 1000 kg1/mole. wt = 700—800. On further heating to viscous liquids with mol. wt = 700—800. Orig. art. has: 3 figures. solid polyesters with mol. wt = 2000. Orig. 678.649:12678.85	
Card 1/2	



MOSHKIMA, T.M.; PUDOVIK, A.N.; ZIL'BERMAN, L.V.

Phosphorus-containing hydrazo and azo compounds. Dokl. AN SSSR 163 no.6:1401-1403 Ag 165. (MIRA 18:8)

- 1. Kazanskiy gosudarstvennyy universitet im. V.I.Ul'yanova-Lenina.
- 2. Chlen-korrespondent AN SSSR (for Pudovik).

Peactions of esters of trivalent phosphorus solds with tisters of pyruvic acid. Zhur. cb. khim. 35 no.9:1501-1505 S 165.

(NUEA 18:10)

1. Kacanskiy gosudarstvennyy universitet.

L 1580-66 EWT(m)/EPF(c)/EWP(j)/T UR/0190/65/007/009/1539/1542 ACCESSION NR: AP5022598 66.095.26+678.86 Pudovik, A. N.; Kuzovleva, R. G. AUTHORS: TITLE: Polymerization and copolymerization of α- and β-carbalkoxyvinyl phosphinates Vysokomolekulyarnyye soyedineniya, v. 7, no. 9, 1965, 1539-1542 SOURCE: TOPIC TAGS: polymerization, copolymer, phosphinate, vinylphosphinate ABSTRACT: Synthesis and polymerization of diethyl  $\ll$ - and  $\beta$ -carbalkoxyvinyl phosphinate (I and II, respectively) have been investigated as a continuation of the study of polymerization and copolymerization of various derivatives of vinylphosphinates reported earlier by the authors (Vysokomolek. soyed., 6, 737, 1964). The two methods for the preparation of I, reported in the literature by J. B. Dickey and H. W. Coover (U. S. Pat. 2559854, Chem. Abstr. 45, 8810, 1951) and A. Ya. Yakubovich, L. Z. Soborovskiy, L. I. Muler, and V. S. Fayermark (Zh. obshch. khimii, 28, 317, 1958) had to be disregarded, as the first one resulted in the wrong product, while the second gave an impure one. Treatment of the diethyl  $\prec$ -chloro- $\beta$ -carbomethoxyethyl phosphinate (from the oxidative phosphorylation Card 1/2

L 1580-66

ACCESSION NR: AP5022598

of methylacrylate) with twice the theoretical amount of triethylamine gave pure I, b.p. 110-112C/2 mm, d<sub>4</sub><sup>20</sup> 1.1687, n<sub>D</sub><sup>20</sup> 1.4389. II, b.p. 131-133C/4 mm, d<sub>4</sub><sup>20</sup> 1.1412, n<sub>D</sub><sup>20</sup> 1.4490 was obtained by using the method of K. M. Kirillova, V. A. Kukhtin, and T. M. Sudakova (Dokl. AN SSSR, 149, 316, 1963). Effects of the temperature, the nature, and the concentration of the initiator on the block polymerization of I and II were studied. It was found that I polymerized to the extent of 80% at 70C after 10 hours in the presence of 1 mole% of benzoyl peroxide. II gave a yield of only 15%. Increase of the temperature to 90C as well as the increase of the concentration of the initiator (azo-bis-isobutyronitrile) resulted in increased yield and polymerization rate. Copolymerization of I with methylmethacrylate, methylacrylate, and acrylonitrile/gave solid polymers. II with methylmethacrylate gave a resinous, soft polymer. Orig. art. has: 1 figure and 1 table.

ASSOCIATION: Kazanskiy gosudarstvennyy universitet im. V. I. Ul'yanova-Lenina (Kazan State University)

SUBMITTED: 120ct64 HH165

ENCL: 00

SUB CODE: OC

NO REF SOV: 004

OTHER: 001

KIRPICHNIKOV, P.A.; MUKMENEVA, N.A.; PUDOVIK, A.N.; YARTSEVA, L.M.

Interaction of a dephenylpicrylhydrazyl with phosphorous acid esters. Zhur. ob.khim. 34 no. 5:1683-1684 My '64.

(MIRA 17:7)

PUDOVIK, A.N.; CHFRKASOV, R.A.

Synthesis of polyalkylene glycol dithiophosphates and some of their addition reactions. Vysokom. soed. 6 no.4:741-744 Ap '64. (MIRA 17:6)

1. Kazanskiy gosudarstvennyy universitet imeni V.I. Ul'yanova-Lenina.

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ACCESSION NR: AT4033992 \$/0000/63/000/000/0091/0095

AUTHOR: Pudovik, A. N.; Cherkasov, R. A.; Pudovik, M. A.

TITLE: Polyalkyleneglycol dithiophosphates and the reactions of their addition to unsaturated compounds

SOURCE: Geterotsepny\*ye vy\*sokomolekulyarny\*ye soyedineniya (Heterochain macro-molecular compounds); sbornik statey. Moscow, Izd-vo "Nauka," 1963, 91-95

TOPIC TAGS: dithiophosphate, polyalkyleneglycol dithiophosphate, polyester, phosphorus containing polyester, polyester synthesis, unsaturated compound, electrophilic unsaturated compound, nucleophilic unsaturated compound, polyester addition reaction

ABSTRACT: Several polyalkyleneglycol dithiophosphates were synthesized by reesterification of dithiophosphoric acid ethers with glycols (ethylene glycol, 1,2-propylene glycol, 1,4-butylene glycol, diethylene glycol, pyrocatechol and hydroquinone). Reactions lasted 1 to 6 hours at 50-130 mm pressure and 80-170C. The resultant polyesters (viscous or nearly solid transparent resins with 15.08 to 20.36% P) were used in additional reactions (30-60 min., 70-80C, 30% excess of the saturated compound, without a catalyst or with sodium ethylate, in dioxane solution for solid or highly viscous polyesters) to acrylonitrile, methacrylate, diethylacrylate, diethylacrylate.

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maleate, styrene, benzylan tables and 4 chemical equa	iline and p-nitrobenzylaniline. tions.	Orig. art.has: 3	
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Card 2/2			

PUDOVIK, A.N.; SITDYKOVA, F.N.

Addition of organophosphorus compounds with a mobile hydrogen to divinyl sulfone and p-di(\$\beta\$-nitrovinyl)benzene. Zhur. ob. khim. 34 no. 5:1682-1683 My '64. (MIRA 17:7)

1. Kazanskiy gosudarstvennyy universitet.

ACCESSION NR: APLIO32576

S/0190/64/006/004/0737/0740

AUTHORS: Pudovik, A. N.; Kuzovleva, R. G.

TITLE: Polymerization and copolymerization of A -acetoxyvinylphosphinic acid esters

SOURCE: Vy\*sokomolek. soyedin., v. 6, no. 4, 1964, 737-740

的现在分词 "你是我们就是我们的现在,我们就是我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我

TOPIC TACS: alkyl vinylphosphinate, methyl vinylphosphinate, ethyl vinylphosphinate, nate, propyl vinylphosphinate, alpha acetoxyvinylphosphinic acid ester, alkyl vinylphosphinate polymerization, alkyl vinylphosphinate copolymerization, methyl methacrylate, methyl acrylate, styrene

ABSTRACT: The polymerization of methyl, ethyl, and n-propyl esters of acetoxyvinylphosphinic acid (AOVPA), and also the copolymerization of these esters with methyl methacrylate, methyl acrylate, and styrene were investigated. The rate of block polymerization was determined by the dilatometric technique, and the yield was estimated by removing the monomer by steam distillation. It was found that in the presence of 2 mole benzoyI peroxide the polymerization rate of AOVPA markedly increased with temperature. At 900 it reached 60% in 10 hours

Card 1/2

ACCESSION NR: AP4032576

(at 50C it took 40 hours to produce 50%). Experiments with various concentrations of benzoyl peroxide at 70C showed an enhancing effect of higher concentrations on the polymerization rate of AOVPA. A comparison of the polymerization rates of methyl, ethyl, and propyl esters of AOVPA revealed that the methyl ester had the lowest polymerization rate, and the propyl ester the highest. Abstracter's note: the authors erred in claiming in the text and conclusions the reverse effect, as evidenced by Chart 27. The copolymerization of AOVPA with methyl methacrylate, methyl acrylate, and styrene was conducted in block, at 70C for a duration of 15 hours, in the presence of 1 mole/% benzoyl peroxide. The copolymers with a small content of AOVPA were hard, transparent products, while the ones containing a larger percentage of AOVPA represented viscous resins. The molecular weights of the copolymers were within the 4580-6616 range, and their vitrification temperature varied from 48C to 87.5C. Orig. art. has: 2 charts and 1 table.

ASSOCIATION: Kazanskiy gosudarstvenny\*y universitet im. V. I. Ul'yanova-Lenina (Kazan' State University)

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Card 2/2\_

TITIE: Reactions of nucleophilic addition to unsaturated polyesters  SOURCE: Vysokomolekulyarnyye soyedineniya, v. 6, no. 12, 1964, 2145-2148  TOPIC TAGS: polyester plastic, macromolecular chemistry  Abstract: Polyesters produced on the basis of maleic anhydride and diethylene glycol. Avith molecular rejects for 2001-1 2004-1	18 11 8
TITIE: Reactions of nucleophilic addition to unsaturated polyesters  SOURCE: Vysokomolekulyarnyye soyedineniya, v. 6, no. 12, 1964, 2145-2148  TOPIC TAGS: polyester plastic, macromolecular chemistry  Abstract: Polyesters produced on the basis of maleic anhydride and diethylene glycol. Avith molecular rejects for 2001-1 2004-1	B
SOURCE: Vysokomolekulyarnyye soyedineniya, v. 6, no. 12, 1964, 2145-2148  TOPIC TAGS: polyester plastic, macromolecular chemistry  Abstract: Polyesters produced on the basis of maleic anhydride and diethylene glycol. A with molecular weights.	
TOPIC TAGS: polyester plastic, macromolecular chemistry  Abstract: Polyesters produced on the basis of maleic anhydride and diethylene glycol. With molecular reights.	
Abstract: Polyesters produced on the basis of maleic anhydride and diethylene glycol. With molecular weights.	
Abstract: Polyesters produced on the basis of maleic anhydride and diethylene glycol. With molecular weights.	
gated in nucleophilic addition reactions. Dialkylphosphorous acids, containing from two to seven carbon atoms in the ester radicals, diethylphosphoneacetic ester, and diethylphosphoneacetone, as well as ammonia and amines (diethylamine and aniline) and malonic, acetoacetic, and cyanoacetic esters were added to the unsaturated polyesters. It was found that these reagents can add to polydiethylene glycol fumerate in the presence of alkali metal alcoholates, and in certain cases (amines) in the absence of catalysts as well. The properties of the resultant products were investigated. It was also shown that electrophilic reagents, in particular,	c

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corresponding to polydiet	unsaturated polyesters; and hylene glycol dichlorosucci	nate was optained when
fluorine was passed throu	gh a cooled chloroform solu	tion of polydiethylene
glycol fumarate with mole	cular weight 894. Orig. art.	has 2 formulas and 3 tables.
ASSOCIATION: Kazanskiy gos	udarstvennyy universitet im.	V. I. Ul'yanova-Lenina
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SJEMITTED: 29 Jan64 No REF SOV: 001	ENCL: 00 OTHER: 001	SUB CODE: MT, GC JPRS

PUDOVIK, A.N.; RAKOV, A.P.

Thermal rearrangement of complete phosphorous acid esters with saturated radicals. Dokl. AN SSSR 161 no.6:1352-1355 Ap 165.

(MIRA 18:5)

1. Kazanskiy gosudarstvennyy universitet im. V.I.Ul'yanova-Lenina.

2. Chlen-korrespondent AN SSSR (for Pudovik).

ACCESSION NR. AT4033991

s/0000/63/000/000/0087/0090

AUTHOR: Nikitina, V. I.; Maklakov, A. I.; Balakireva, R. S.; Pudovik, A. N.

TITLE: Polymers consisting of aromatic rings conjugated with hetero atoms. I. Polypheneylene- and polydiphenyleneimines.

SOURCE: Geterotsepny\*ye vy\*sokomolekulyarny\*ye soyedineniya (Heterochain macromolecular compounds); sbornik statey. Hoscow, Izd-vo "Nauka," 1963, 87-90

TOPIC TAGS: organic semiconductor, semiconducting polymer, polyphenyleneimine, polydiphenyleneimine, polymer electrical property

ABSTRACT: Polyphenyleneimines, semiconducting polymers containing
NH groups between aromatic rings in the backbone, have been prepared,
and their electrical and magnetic properties have been studied at the
Kazanskiy gosudarstvenny\*y universitet im. V. I. Ul'yanov-Lenina
(Kazan' State University). Polycondensation of aromatic diamines with
dihydric phenols was used. Polymer I was obtained from p-phenylenediamine and hydroquinone; II, from bensidine and hydroquinone;

# ACCESSION NR. AT4033991

and III, from benzidine and 4,4'-dihydroxybiphenyl. The polycondensation was carried out in the melt at 260—280C for 6 hr, and then at the same temperature and 1—2 mm Hg for 2 hr. All the polymers are black powders insoluble in the common organic solvents except dimethylformamide. Three types of polymers were prepared: repractipitated (a), nonreprecipitated (b), and reprecipitated and repractipitated in air at 320C for 1 hr (c). On the basis of the synthesis method and IR spectroscopy, the following structures were

$$-\left[-NH-\right]^{n}-\left[-NH-\right]^{n}$$
(III)

assumed. D-c electrical conductivity was determined for pellet samples at 20—100C. The temperature dependence of conductivity obeyed an exponential law fairly well. Numerical data are given in Table 1 of the Enclosure. All the samples exhibited high conductivity, positive magnetic susceptibility, and an EPR signal. The unpaired

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			T	able 1.	Ele	ctrica molyp	l and ma nenylene	gneti imin <b>e</b> :	c prope	rties		2	
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		I	a * b c	10 <sup>-9</sup> 10 <sup>-10</sup> 10 <sup>-11</sup>	10 <sup>-4</sup> 10 <sup>-4</sup> 10 <sup>-6</sup>	0.48 0.60 0.68	~7	111	10 <sup>-9</sup> 10 <sup>-12</sup> 10 <sup>-10</sup>	10-5	0.68 0.86 0.50		
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PUDOVIK, A.N.; KUZOVLEVA, R.G.

Polymerization and copolymerization of  $\alpha$  -acetoxyvinylphosphinic Polymerization and copolymerization acid esters. Vysokom. soed. 6 no.4:737-740 Ap 164. (MIRA 17:6)

1. Kazanskiy gosudarstvennyy universitet imeni V.I. Ul'yanova-Lenina.

L 21810-65 EPA(s)-2/EWT(m)/EPF(c)/EPR/EWP(j)/T Pc-4/Pr-4/Ps-4/Pt-10 WW/RM ACCESSION NR: AP5001478 S/0190/64/006/010/2139/2144

AUTHOR: Pudovik, A. N.; Yevstaf yev, G. I.

TITLE: Synthesis of polyphosphinites by transesterification

SOURCE: Vysokomolekulyarnyye soyedineniya, v. 6, no. 12, 1964, 2139-2144

TOPIC TAGS: phosphorus containing polymer, polyphosphinite, polyglycolphosphinite, glycol, polyethylene glycol, self extinguishing polymer, transesterification

ABSTRACT: As a part of their study of the formation of phosphorus(III) containing polyesters, the authors obtained poly(alkylene glycol phenylphosphinites) and poly(alkylene glycol ethylphosphinites) by the transesterification of dialkyl phenyl- or dialkyl ethyl-phosphinite. It was demonstrated that the polycondensation of these esters with aliphatic glycols, such as ethylene, diethylene, or tetraethylene glycol, essentially takes place via cyclic phosphinites. These cyclic esters seem to be quite stable, even when the number of atoms in the

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ACCESSION NR: AP5001478

ring exceeds six. Pure cyclic esters, isolated in the study, are liquids with a phosphorus content close to the theoretical. On standing, they polymerize to oligomers; on heating over 140C, high-molecular-weight products insoluble or slightly soluble in benzene or dioxane are formed. The polyesters obtained are viscous liquids; when burned, some of them are self-extinguishing after being taken out of the flame. Cyclic glycol phosphinites and their polymers contain trivalent phosphorus; this is indicated by addition reactions, e.g. with sulfur, in which phosphonothicate derivatives are formed. Orig. art. has: 1 formula, 4 figures, and 2 tables.

ASSOCIATION: Kazanskiy gosudarstvennyy universitet im. V. I. Uliyanova-Lenina (Kazan State University)

SUBMITTED: 29Jan64

ENCL: 00

SUB CODE: OC. HT

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